

A stainless steel strip coated with a decorative layer

The present invention relates to a new coated stainless steel strip material with a covering layer with a decorative surface finish. It also relates to a method of 5 manufacturing such a coated steel strip in a continuous roll-to-roll process, which results in a very good adhesion of a decorative covering layer on a metal strip substrate. In particular, it relates to coated steel strips with good formability and with such a good adhesion of the decorative 10 layer that they are suitable for use in consumer related applications, although many other uses also are feasible.

Background to the Invention and Prior Art

It is known that decorative coatings can give an 15 attractive surface finish to consumer related products. However, for components in smaller dimensions, which are to be produced in a cost-efficient and productive way, there are difficulties in finding a method that can attain the quality and productivity requirements. For productivity 20 reasons, a roll-to-roll coating process is imperative, and for quality reasons, a thin layer with excellent adhesion is needed.

The superior adhesion is required for the functional quality of the final product, but also to enable a cost-efficient and productive manufacturing of components. Thus, 25 a coated strip material with inferior adhesion would cause problems with, e.g., flaking, and this would result in a low yield and also in a disturbance caused by the flakes themselves of the manufacturing process as such, especially if the manufacturing process is in a continuous line. Moreover, more frequent stops would be needed for quality 30 inspections and for cleaning the process line from flakes.

All in all, poor adhesion of the coating would result in a non-acceptable high manufacturing cost and low quality.

There are several common methods of making a decorative surface finish on metallic materials. As examples can be mentioned:

■ Anodizing is a known method that can be used for various colours. Normally this method is used on aluminum or aluminum alloys. An obvious drawback is that it is not possible to use directly on stainless steels.

10 ■ Vapor deposition methods are used in some cases for colouring of metallic products. Often the colour is produced by applying a metal nitride to the surface of the component. However, most methods are batch-like processes, which mean that the coating is done on the finished component piece by piece. One obvious drawback with such a method is that it is not continuous, and thus also very expensive to use. Examples of batch coating on consumer related products are revealed in US 6 197 438 B1, US 5 510 012, and EP-A-1 033 416.

20 ■ Recent developments in PVD technology has led to the manufacturing of continuous PVD coating lines for various reasons, so called web-coaters are commonly used to coat plastic foils for food packing i.e. chips bags etc. Continuous PVD lines are also used for the 25 manufacturing of highly reflective surfaces on aluminum strips, semi-continuous PVD lines are also commonly used to coat window glass. Less common are continuous PVD lines for coating stainless steel materials with functional layers. In US 6 197 132 a continuous PVD line 30 for coating stainless steel strip with aluminum is described. Also in the article "Innovative steel strip coatings by means of PVD in a continuous pilot line: process technology and coating development" by B. Schuhmacher et al., Surface and Technology 163-164,

(2003) pp 703-709, a continuous PVD process for depositing Zn-alloys as corrosion protection.

Furthermore, in US 4 763 601 a continuous line for coating of steel strip is described. In said coating

5 line several different coating zones are integrated, including ion plating, sputtering and plasma CVD.

However, the strip speeds used in US 4 763 601 would render a very low productivity. Moreover, the low feed speed of the belt causes long presence times in the
10 coating zones which could result in a deteriorated end product.

- One commonly used method is painting of the metallic surface with coloured lacquers, or the like. However, in most painting processes the painting is done on the
15 finished component piece by piece. One obvious drawback with such a method is that it is not a continuous roll-to-roll process, and thus also quite expensive to use. A continuous painting process is normally not possible to use since the adhesion usually is not good enough for
20 further processing in, e.g., forming operations without causing defects or flaking in the surface. Also, paints can normally not withstand further heat treatments.

Thus, the methods as described in the examples above cannot be used for the present invention.

25 Therefore, it is a primary object of the present invention to provide a decorative coated metal strip with good adhesion between the decorative coating and the substrate.

A further object of the present invention is to obtain a
30 cost-efficient decorative coating in a continuous roll-to-roll process with a minimum strip speed of 3m/min, integrated in the production of a stainless steel strip.

Yet another object of the present invention is to provide a coated stainless steel strip product with a decorative coating in combination with good formability, so as to enable the manufacturing of consumer related applications of said material.

Furthermore, an object of the present invention is to provide a coated stainless steel strip with a decorative layer that if desired can be heat-treated in a tempering or hardening process without changing its colour.

Still another object of the present invention is to provide a method to manufacture a surface finish in connection to a continuous coating in a roll-to-roll process included in a strip production line, which in the further processing by a suitable heat-treatment, can be transformed to a decorative coating.

A further object of the present invention is to obtain a coating with a thickness as uniform as possible.

These and other objects have been attained by providing a coated steel product according to claim 1. Further preferred embodiments are defined in the dependent claims.

Brief Description of the Invention

The present invention relates to a method of manufacturing coated stainless steel in a continuous roll-to-roll process, which results in an excellent adhesion of a thin covering decorative layer. The decorative layer can as an example be achieved by applying a coating so as a colour appears. The coated stainless steel strips must have such a good adhesion of the thin layer that it is suitable for a cost-efficient and productive manufacturing of components in consumer related applications. The final product, in the form of a decorative stainless strip material, is suitable for use as a decorative component in

consumer-related applications that are occasionally used also in environments with high humidity or in conditions requiring a good corrosion resistance.

The decorative layer is deposited by means of Physical Vapor Deposition (PVD) in a roll-to-roll process, to an evenly distributed layer with a thickness of preferably less than 10 µm. The preferred PVD methods to be used are either electron beam evaporation (EB) or sputtering and well known to the skilled man.

The substrate material can preferably be a stainless steel with a Cr content above 10% (by weight) and with a strip thickness of usually less than 3 mm. As a first step, the roll-to-roll process may also include an etch chamber, in order to remove the native oxide layer that otherwise normally is present on a stainless steel.

As an alternative, the decorative surface finish may also be produced in a two-step process, in which first a suitable coating is applied to the stainless steel strip material in a condition allowing good formability, using a method as described above, and thereafter in a subsequent processing of the already coated material, use a suitable heat-treatment, in which the thin coated layer transforms to a decorative surface finish.

25 Brief Description of the Drawings

Figure 1 shows an illustration of a test specimen in accordance with the present invention, i.e. a coated stainless steel strip with a decorative layer with good adhesion before a test of said adhesion in a bend test over a radius maximally equal to $5^* t$, where t is the thickness of said strip.

Figure 2 shows an illustration of a test specimen in accordance with the present invention, i.e. a coated stainless steel strip with a thin decorative layer with

good adhesion, and after a bending in an angle of 180°, in a bend test as described in Fig.1.

Figure 3 shows schematically an electron beam evaporation production line for the manufacturing of a 5 coated metal strip material according to the invention.

Figure 4 shows schematically a sputtering production line for the manufacturing of a coated metal strip material according to the invention.

Figure 5 shows an illustration of the CIE lab 10 ordinates L*, a* and b* values, in which the L* value is the brightness from black to white, the a* value goes from green to red and the b* value is blue to yellow.

15 Detailed Description of the Invention

Description of the Coating and the Use of the Invention

The final product, in the form of a coated strip material, is suitable to be used as a decorative component in consumer-related applications such as outdoor life 20 applications, sports and sea-life applications, household applications, camera applications, mobile phones and other telecom applications, edge applications such as knife, saw and shaving applications or the like, and applications for personal belongings and care such as watches, glasses, 25 cosmetic applications, buttons and zippers in clothing, perfume bottles or the like. In principle, these are all applications that preferably can be given an attractive surface finish that by being decorative gives an added design value to the final product. Also, they are 30 occasionally used in environments with high humidity or in conditions requiring good corrosion resistance. At the same time, these types of applications are often expected to be nice-looking throughout its product lifetime, with a shiny appearance, or just a "high quality" appearance. Dull

surfaces, with spots or even rust, are normally not acceptable.

Both one-sided and two-sided coatings may be used, but since the component normally is used in the final consumer related product in such a way that only one side is visible, i.e., the external surface, it is preferable from a cost perspective to use one-sided coatings whenever possible. A substrate material in stainless steel also makes this possible, since the coating does not have to act as a corrosion protection in itself, and thus from a corrosion point of view, it does not have to cover the component on all sides.

The method described in the present invention is especially suitable for thin coatings in thicknesses on each side up to 10 μm in total, normally up to 7 μm in total, preferably up to 5 μm in total, or at the best maximum 3 μm , or even maximum 2 μm in total, is preferable from a cost perspective. If thicker layers are to be coated, an optimum in cost versus properties may be achieved by using multi-layers with up to 10 layers, and where each layer is between 0,01 to 10 μm thick, suitably between 0,01 to 5 μm , preferably 0,01 to 3 μm and even more preferably 0,005 to 2 μm .

The thickness tolerances obtained by PVD techniques are usually very good. Thus, the tolerances of each layer may be maximally +/- 30% of the layer thickness in strip widths up to 400 mm, normally +/- 20%, and preferably +/- 10%, most preferably +/- 7%. This means that very tight tolerances can be achieved, which is of benefit for the precision during usage and the quality of the product. A tight tolerance in layer thickness is also of advantage for achieving a consistency in colour. Thus, thanks to the high coating thickness tolerances, a superior colour consistency has been achieved, even on long belts, such as 5 km and

even longer. These long belts have been made possible thanks to the relatively high feed velocities. Colour differences expressed as ΔE (see definition further down) of less than 10 have been achieved for those long belts.

- 5 For shorter belts, such as up to 100 m ΔE may even be less than 5.

Moreover, thanks to the fact that the belt may be cooled by special cooling means at the same time as it is coated, the cooling taking place on the side of the belt 10 opposed to the side being coated, the heat influence on the belt is largely controlled, so that the properties of the substrate are substantially maintained.

The thin covering layer must have a good adhesion in order for the coated strip product to have a good 15 formability during manufacturing to various components, not to cause problems due to flaking of the layer or the like. The thin layer must also have a good adhesion with regard to the applications and their uses. During usage it is not acceptable that the decorative layer starts to flake off. 20 An illustration of the good adhesion is that with the substrate material in a soft annealed condition, the coated stainless steel strip according to the present invention should be able to be bent a certain minimum angle over a radius maximally $5 * t$, where t is the thickness of said strip, without showing any tendency to flaking or the like. 25 The minimum angle in this type of bending is typically minimum 90° , normally minimum 120° , preferably minimum 150° and even more preferably 180° (See Fig 1-2). According to one embodiment of the invention the coated strip should be 30 able to be bent at least 90° , preferably at least 180° , over a radius that is equal to the thickness of the strip without showing any tendency to flaking or the like.

The coated layer should be sufficiently wear-resistant in order to withstand the wear and shear exerted by the

treated material, on the other hand it should not be too thick, due to economical reasons and fragility/brittleness. For consumer related applications, the ratio between the thickness of the coating and the substrate material should 5 be between 0,001% to 7 %, normally 0,001 to 5 % and usually 0,001 to 4 %, but most preferably between 0,001-3 %.

The decorative appearance can be achieved by depositing several different compounds or elements. Metal oxides are well known to have a decorative appearance and a 10 suitable decorative coating according to the present invention is to use coatings of binary metal oxides such as SiO_2 , SiO , TiO , VO , VO_2 , TiO_2 , Al_2O_3 , Cr_2O_3 , CoO , NiO , Cu_2O , CuO and as well as coatings of ternary metal oxides such as BaTiO_3 , PbCrO_4 , Pb_2CrO_5 , NaWO_3 , NaWO_2 , MgAl_2O_4 , TiMg_2O_4 , 15 LaCrO_3 are decorative. In addition solid solutions and/or mixtures of metal oxides can also be deposited to achieve decorative coatings on the strip substrates. These mixtures or solutions of metal oxides are preferably, Al_2O_3 , TiO_2 or Cr_2O_3 based systems. A ternary metal compound 20 such as a ternary oxide can be deposited by means of co-evaporation or by sputtering.

In variation to the above described coatings, also deposited layers of metal nitrides such as TiN , ZrN , CrN , TiAlN are all good examples of a good way to achieve 25 decorative coatings. Also metal carbides and metal carbo-nitrides such as TiC , TiCN are coatings, which have a decorative appearance. By using this type of coating, the coated strip material can, if desired, be subsequently heat-treated in a suitable gas atmosphere of normally for 30 metal nitrides a nitrogen containing gas, such as N_2 , NH_3 , N_2H_4 or mixtures thereof, or for metal carbides a carbon containing atmosphere such as CH_4 , CH_3CH_3 , C_2H_4 , C_2H_2 or mixtures thereof, or for metal carbonitrides a gas consisting of a mixture of nitrogen containing and carbon 35 containing gases. The decorative layer can after such a

heat-treatment keep its colour achieved in the coating as expressed in terms of L*, a*, b*-values, in a way that the difference in decorative appearance, or colour, between the layer before the heat-treatment and after the heat-treatment as expressed in terms of ΔE, is in strip widths up to 400 mm maximally 15, usually maximally 12, normally 8, preferably 5 and even more preferably 3, and at the best even 1.

This type of coating is therefore of special interest to use for consumer related applications requiring also a high mechanical strength, such as edge applications, e.g. knife applications, saw applications or shaving applications such as a razor blade, or other high-strength applications, e.g. out-door applications, glasses or perfume bottles. This is all applications typically manufactured in materials such as hardenable chromium steels, precipitation hardenable stainless steels or cold-rolled stainless spring steels, which normally are heat-treated in a tempering process effected at temperatures in the range 325–525 °C, and preferably in between 375–500 °C, or in a hardening done at high temperatures, usually above 400 °C, normally above 800 °C and in some cases above 950 °C. After such a heat-treatment as described here, the substrate material increases its mechanical (tensile) strength to usually above 1000 MPa, typically above 1200 MPa and normally above 1400 MPa, and preferably even above 1500 MPa. This heat-treatment can be performed on the coated strip material in a continuous roll-to-roll process, or on components manufactured from the coated strip product. In the latter case the second step, e.g. the heat-treatment, is done as a piece by piece process.

Multi-layers may also be used in order to enable a combination of oxides and/or nitrides or carbides, so as to optimize colour spectra, reflectiveness and consistency

in colour by having up to 10 layers with different oxides and/or nitrides or carbides in the layers.

There is normally no need for any separate interlayer bond-coat, but it may still be used in at least one of the 5 layers if it is required from a technical perspective, e.g., to enhance toughness. A suitable bond-coat to use is a pure metal such as nickel, aluminum, chromium, titanium or the like. Since a separate metallic bond-coat layer means an extra cost it is usually used in very thin layers 10 only, suitably between 0 to 2 μm , preferably between 0-1 μm and most preferably between 0-0,5 μm . The interlayer bond-coat may also be used in between the substrate material and the first layer of decorative coating, especially if the first layer is made of a non-metallic coating.

15 An alternative method to the above described, is to use a two-step method to produce the decorative surface finish. In the first step a suitable coating is applied to the steel strip material using a method as described in the present invention. In this case a suitable coating is a 20 covering layer of a metal such as aluminum, chromium, titanium, zirconium, vanadium, hafnium, copper, nickel, cobalt or a binary oxide of said metals such as Al_2O_3 , Cr_2O_3 , but it can also be mixtures of these metals and their corresponding metal oxides. In this case a specific 25 thickness and composition of the coating is the most important in the first step, and not the colour itself. The coated steel product should in this stage also be of good formability and have a good adhesion of the substrate and the thin layer, so as to be able to be manufactured in 30 various forming processes as described in the present invention, and as illustrated by the bend-test as described in Fig 1-2. In a subsequent processing by a selected chemical treatment and/or heat-treatment, the layer can be transformed in composition so that a desired colour is

formed on the surface of the material. Preferably, this subsequent treatment is made in connection to an ordinary process, such as during a tempering treatment of a cold rolled spring steel or a precipitation hardenable steel or 5 during a hardening operation of a hardenable chromium steel, or the like. Tempering treatments as described above may be utilised. By selecting an atmosphere in the furnace to be reactive with regard to the coating, the desired transformation of the coating can be achieved.

10 If it is desirable to have on the final coated strip an oxide coating, then the coated metal layer and/or binary oxide is heat treated in an oxidizing atmosphere, the oxidizing atmosphere can be water vapor, pure oxygen gas, O₂, or different mixtures of oxygen with an inert gas such 15 as Ar, N₂, He, or at the best air at a suitable temperature.

If the desired is to incorporate nitrogen into the metal and/or metal oxide coating, then a reactive gas containing nitrogen may be used during the heat-treatment, including 20 gases such as N₂, NH₃, N₂H₄ or mixed gases of N₂/H₂. The final decorative layer is after such a heat treatment transformed into either a metal nitride or a metal oxynitride.

25 If the desired is to incorporate carbon into the metal and/or metal oxide coating, then a reactive gas containing carbon may be used as reactive atmosphere during the heat-treatment that will include gases such as CH₄, CH₃CH₃, C₂H₄, C₂H₂ or mixed gases of these. If the final decorative layer 30 should be metal carbo-nitride, the carbon containing gases are mixed with nitrogen containing gases and heat-treated at elevated temperatures.

By using this type of two-step process, an additive element such as oxygen, nitrogen, carbon have been incorporated into the as-deposited layer transforming it 35 into a metal oxide, metal nitride, metal carbide or a mixed

type of layer, usually in the form of an oxy-nitride, oxy-carbide or carbo-nitride resulting in a final product with an attractive decorative colour of the surface. The subsequent processing as described here can be performed on
5 the coated strip material in a continuous roll-to-roll process, or on components manufactured from the coated strip product. In the latter case the second step, e.g. the heat-treatment, is performed as a piece by piece process.

The final product in the form of a coated strip
10 material in accordance with the present invention should also be capable of being readily manufactured to components suitable for applications as described above, in a cost-efficient and productive manufacturing process, including forming steps such as deep-drawing, punching, stamping, or
15 the like. [cf. Figures 1 and 2]

Description of the decorative surface finish

Colour is the way the HVS (the human visual system) measures a part of the electromagnetic spectrum,
20 approximately between the wave-lengths 300-830 nm. Because of certain properties of the HVS we are not able to see all of the possible combinations of the visible spectrum but we tend to group various spectra into colours. A colour space is a notation by which we can specify colours i.e. the
25 human precipitation of the visible electromagnetic spectrum.

CIE, the International Commission on Illumination-abbreviated as CIE from its French title Commission Internationale d'Éclairage - is an organisation devoted to
30 international co-operation and exchange of information among its members countries on all matters relating to science and art of lighting.

CIE standardised the XYZ values as tristimulus values that describe any colour that can be perceived by an

average human observer. These primaries are nonreal, i.e. they cannot be realised by actual colour stimuli. This colour space is chosen in such a way that every perceptible visual stimulus be described with positive **XYZ** values. A very important attribute of the CIE **XYZ** colour space is that it is device independent.

The transformation from CIE **XYZ** to CIE Lab is performed with following equations

$$10 \quad L^* = 116 \left(\frac{Y}{Y_n} \right)^{\frac{1}{3}} - 16$$

$$a^* = 500 \left[\left(\frac{X}{X_n} \right)^{\frac{1}{3}} - \left(\frac{Y}{Y_n} \right)^{\frac{1}{3}} \right]$$

$$b^* = 200 \left[\left(\frac{Y}{Y_n} \right)^{\frac{1}{3}} - \left(\frac{Z}{Z_n} \right)^{\frac{1}{3}} \right]$$

15 The trimulus values **Xn**, **Yn**, **Zn** are those of the normally white objective-colours stimulus. The **L*** value is the brightness from black to white, the **a*** value goes from green to red and the **b*** value is blue to yellow, see also Figure 5.

20 The perceptually linear colour difference formulas between two colours is defined as:

$$\Delta E = \sqrt{(\Delta L)^2 + (\Delta a)^2 + (\Delta b)^2}$$

25 The colour can be describe in **L***, **a*** and **b*** value.

The decorative coatings in the present invention is suitable for a colour space in following parameter of **L***,

a^* and b^* . The parameter of L^* , a^* and b^* are usually $0 < L^* < 95$, $-66 < a^* < 64$ and $-90 < b^* < 70$. A blue colour is normally $20 < L^* < 95$, $-66 < a^* < 64$ and $-83 < b^* < 0$. A green colour is normally $20 < L^* < 95$, $-66 < a^* < 0$ and $-83 < b^* < 70$.

5 A red colour is normally $20 < L^* < 95$, $0 < a^* < 64$ and $-40 < b^* < 35$. A golden colour is normally $20 < L^* < 95$, $-66 < a^* < 64$ and $0 < b^* < 70$. A black colour is normally $0 < L^* < 50$, $-20 < a^* < 20$ and $-20 < b^* < -20$. A violet colour is normally $20 < L^* < 95$, $20 < a^* < 60$ and $-25 < b^* < -60$.

10 A blue colour is preferably $20 < L^* < 50$, $-15 < a^* < 50$ and $-70 < b^* < 0$. A green colour is preferably $60 < L^* < 90$, $-60 < a^* < -7$ and $-83 < b^* < 70$. A red colour is preferably $40 < L^* < 60$, $20 < a^* < 40$ and $-10 < b^* < 10$. A golden colour is preferably $60 < L^* < 95$, $-10 < a^* < 25$ and $30 < b^* < 50$. A black colour is preferably $0 < L^* < 40$, $-10 < a^* < 10$ and $-10 < b^* < 10$. A violet colour is preferably $20 < L^* < 50$, $20 < a^* < 60$ and $-25 < b^* < -60$.

15 The method described in the present invention is especially suitable to use when a good consistency in the colour is desired.

Description of the substrate material to be coated

The material to be coated should have a good basic corrosion resistance, preferably with a chromium content of more than 12%, or at least 11% or minimum 10% (by weight), depending on the composition of the other alloying elements. It should also be possible to use in a state of good formability, which in a soft-annealed condition may be expressed in terms of elongation measured as A50 of more than 1%, normally more than 2%, and preferably more than 3%. Materials that are suitable to use are alloys such as ferritic chromium steels of the type AISI 400-series,

austenitic stainless steels of the type AISI 300-series, hardenable chromium steels, duplex stainless steels, or precipitation hardenable stainless steels, such as the alloy disclosed in WO 93/07303. This means usually a composition of essentially (wt %):

- Ferritic stainless steel, or an Austenitic stainless steel, or a duplex stainless steel of:
0,001-0,7% C, 10-26% Cr, 0,01-8% Mn, 0,01-2% Si, 0,001-16%
10 Ni, up to 6% Mo, 0,001-0,5% N, up to 1,5% Al, up to 2% Cu,
Nb+W+V in total up to 1% and rest essentially Fe, or:
- Hardenable chromium steels of 0,1-1,5% C, 10-16% Cr,
0,001-1% Ni, 0,01-1,5% Mn, 0,01-1,5% Si, up to 3% Mo,
0,001-0,5% N, Nb+W+V in total up to 1% and rest essentially
15 Fe; or
- Precipitation hardenable steels of: 0,001-0,3% C, 10-16%
Cr, 4-12% Ni, 0,1-1,5% Ti, 0,01-1,0% Al, 0,01-6% Mo, 0,001-
4% Cu, 0,001-0,3% N, 0,01-1,5% Mn, 0,01-1,5% Si, Nb+W+V+Ta
in total up to 2%, rest essentially Fe.

20 Also other stainless grades such as Cobalt alloyed steels,
high-Ni alloys or Ni-based alloys may also be used. The
substrate material can also be in different conditions,
depending on the requirement of mechanical properties, such
as in soft annealed or cold-rolled condition or even
25 hardened condition. The substrate material should in soft-
annealed condition have a tensile strength of maximum 1400
MPa, preferably max 1000 MPa, in order to have a good
formability. In cold-rolled condition should the substrate
material have a tensile strength of minimum 500 MPa,
30 normally min 700 MPa and preferably min 1000 MPa, and in
hardened condition a minimum tensile strength of usually
1000 MPa, or more normally min 1200 MPa. The coating method
may be applied on any kind of product made of said type of
stainless steel alloys and in the form of strip, bar, wire,
35 tube, foil, fiber etc., preferably in the form of strip or

foil, that have good hot workability and also can be cold-rolled to thin dimensions. The alloy should also readily be manufactured to components in a productive manufacturing process including steps such as forming, deep drawing,

5 punching, stamping, or the like.

The thickness of the strip substrate material is usually between 0,015 mm to 3,0 mm and suitably between 0,03 mm to 2 mm. Preferably; it is between 0,03 to 1,5 mm, and even more preferably between 0,03 to 1 mm. The width of
10 the substrate material depends on if the coating is made before or after any foreseen slitting operation. Further, said width should preferably be selected to be a width suitable for further manufacturing to the final width of the component intended to be used in a consumer related
15 application. In principle, the width of the substrate material is therefore between 1 to 1500 mm, suitably 1 to 1000 mm, or preferably 1 to 500 mm, or even more preferably between 5 and 500 mm. The length of the substrate material is suitably at least 10 m, normally between 50 and 20 000 m
20 and preferably between 100 and 20 000 m. According to an embodiment the substrate material is at least 500 m long.

Description of the Coating Method

A variety of physical or chemical evaporation deposition methods for the application of the coating media and the coating process may be used as long as they provide a continuous uniform and adherent layer. As exemplary of deposition methods can be mentioned chemical vapor deposition (CVD), metal organic chemical vapor deposition
25 (MOCVD), physical vapor deposition (PVD) such as sputtering and evaporation by resistive heating, by electron beam, by induction, by arc resistance or by laser deposition methods, but for the present invention especially two PVD methods are preferred for the deposition, either electron
30 beam evaporation (EB) or sputtering. Optionally, the EB
35

evaporation can be plasma activated to even further ensure good quality coatings of dense and decorative layers.

For the present invention, it is a pre-requisite that the coating method is integrated in a roll-to-roll strip production line with a minimum strip speed of 3 m/min to achieve a cost efficient productivity and also to be able to maintain the properties of the substrate material by minimizing the heat influence, which otherwise would risk to deteriorate the properties of the end-product. The coating layer is then deposited by means of electron beam evaporation (EB) or by sputtering in a roll-to-roll process. If multi layers are needed; the formation of them can be achieved by integrating several EB deposition chambers or sputtering chambers in-line. The deposition of metallic coatings should be made under reduced atmosphere at a maximum pressure of 1×10^{-2} mbar with no addition of any reactive gas to ensure essentially pure metal films. The deposition of metal oxides should be performed under reduced pressure with an addition of an oxygen source as reactive gas in the chamber. A partial pressure of oxygen should be in the range $1 - 100 \times 10^{-4}$ mbar. If other types of coatings are to be achieved, e.g., metal carbides and/or nitrides such as TiN, TiC or CrN, or mixtures thereof the conditions during the coating should be adjusted with regard to the partial pressure of a reactive gas so as to enable the formation of the intended compound. In the case of oxygen, a reactive gas such as H₂O, O₂ or O₃, but preferably O₂, may be used. In the case of nitrogen a reactive gas such as N₂, NH₃ or N₂H₄, but preferably N₂, may be used. In the case of carbon, any carbon containing gas may be used as reactive gas, for an example CH₄, C₂H₂ or C₂H₄. All these reactive EB evaporation processes may be plasma activated.

To enable a good adhesion, different types of cleaning steps are used. First of all, the surface of the substrate

material should be cleaned in a proper way to remove all oil residues, which otherwise may negatively affect the efficiency of the coating process and the adhesion and quality of the coating layer. Moreover, the very thin
5 native oxide layer that normally always is present on a steel surface must be removed. This can preferably be done by including a pre-treatment of the surface before the deposition of the coating. In this roll-to-roll production line, the first production step is therefore preferably an
10 ion assisted etching of the metallic strip surface to achieve good adhesion of the first covering layer [see Fig. 3].

Description of embodiments of the invention

15 Six examples of embodiments of the invention will now be described in more detail.

Firstly, the substrate materials are produced by ordinary metallurgical steel making to a chemical composition as described in the specific examples. They are
20 afterwards hot-rolled down to an intermediate size, and thereafter cold-rolled in a conventional manner in several steps with a number of recrystallization steps between said rolling steps, until a final specific thickness of normally < 3 mm, and a width of maximum 400 mm. The surface of the
25 substrate material is then cleaned in a proper way to remove oil residuals from the rolling. Thereafter, the coating process takes place in a continuous process line, starting with decoiling equipment. The first step in the roll-to-roll process line can be a vacuum chamber or an
30 entrance vacuum lock followed by an etch chamber, in which ion-assisted etching takes place in order to remove the thin oxide layer on the surface of the stainless substrate material. The strip then enters into the E-beam evaporation chamber(s) or sputtering chamber(s) in which the deposition

takes place. A layer of normally 0,01 up to 10 μm is deposited in a ratio between the thickness of the coating and the thickness of the strip to be up to max 7%, the preferred thickness depending on the application. In the 5 six examples described here, the specific thickness of the layer is deposited by using one E-beam evaporation chamber and/or one sputtering chamber.

The roll-to-roll PVD process referred to above is illustrated in Figure 3-4. In Figure 3 as an embodiment of 10 the present invention the deposition method is electron beam evaporation and the first part of such a production line is the uncoiler 13 within a vacuum chamber 14, then the in-line ion assisted etching chamber 15, followed by a series of EB evaporation chambers 16, the number of EB 15 evaporation chambers needed can vary from 1 up to 10 chambers, this to achieve a multi-layered structure, if so desired. All the EB evaporation chambers 16 are equipped with EB guns 17 and water-cooled copper crucibles 18 for the evaporation. After these chambers come the exit vacuum 20 chamber 19 and the recoiler 20 for the coated strip material, the recoiler being located within vacuum chamber 19. The vacuum chambers 14 and 19 may also be replaced by an entrance vacuum lock system and an exit vacuum lock system, respectively. In the latter case, the uncoiler 13 25 and the coiler 20 are placed in the open air. After the EB evaporation, the coated strip material passes through the exit vacuum chamber or exit vacuum lock before it is being coiled on to a coiler. In Figure 4 as a further embodiment to the present invention the deposition method is 30 sputtering and the first part of such a production line is the uncoiler 23 within a vacuum chamber 24, then the in-line ion assisted etching chamber 25, followed by a series of sputtering chambers 21, the number of sputtering chambers needed can vary from 1 up to 10 chambers, this to

achieve a multi-layered structure, if so desired. All the sputtering chambers 21 are equipped with dual magnetrons 22 for the deposition. After these chambers comes the exit vacuum chamber 27 and the recoiler 28 for the coated strip 5 material, the recoiler being located within vacuum chamber 27. The vacuum chambers 24 and 27 may also be replaced by an entrance vacuum lock system and an exit vacuum lock system, respectively. In the latter case, the uncoiler 23 and the coiler 28 are placed in the open air. After the 10 sputtering, the coated strip material passes through the exit vacuum chamber or exit vacuum lock before it is being coiled on to a coiler.

As an even further embodiment to the present invention, the deposition method is both electron beam 15 evaporation chambers 16 and sputtering chambers 21 integrated in a series of deposition chambers in a similar production process as described in Figures 3 and 4.

The coated strip material can now, if needed, be further processed by, for example, annealing, rolling or 20 slitting, to obtain the preferred final dimension and condition for the manufacturing of components.

The final product as described in the six examples, i.e., a coated stainless strip material in the specific strip thickness and with a thin covering coating layer in 25 accordance to the examples described below, have a very good adhesion of the coated layer and are thus suitable to be used in a cost-efficient and productive manufacturing of components in consumer related applications. The good adhesion of the layers is further described in Figures 1-2. 30 A substrate material of a stainless steel strip 1 that has been coated with a thin covering layer 2 so as to produce a coated strip product in accordance with the present invention, is put on to a support 4 with a shaped top that has a radius 5 that is maximally $5*t$, where t is the

thickness 3 of said strip. A bend test is then performed in a way that bends said strip up to 180° over the radius 5 and the bending continues until the selected bend angle is attained, and for a bend-angle of 180° until the strip ends 5 meet 6. The substrate material is normally in soft-annealed condition during a bend-test for testing of the adhesion of the coating, since it is the adhesion that is to be tested and not the basic formability of the substrate material. Therefore, the test is perfomed with the coated strip 10 material in a state in which the limitation of formability is in the coating and not in the substrate material. When the bending has been completed in such a bend test, the test specimen is investigated and especially the quality of the layer after bending 7 and the quality of the substrate 15 after bending 8 and the adhesion between said layer and substrate. The test specimens in accordance with the six examples here have all been tested and the results can be seen in Table 1. The different examples in the form of a coated strip test specimen have been tested in a bend-test 20 as described in Fig. 1-2 in a bend angle of between 90° and 180°. Afterwards all tested specimens have been inspected visually with regard to if any tendency of flaking, cracking of the layer, or the like, can be seen. Also, all 25 test-specimens tested in a bend-angle of 180° have been investigated with regard to adhesion with an adhesive Scotch tape, and inspected afterwards if any flakes coming from the coated layer can be seen on the tape. A coated strip material as tested according to this description should have such a good adhesion of the layer that a radius 30 of maximum 5*t, usually 3*t, normally 2*t, preferably 1,5*t, and even more preferably 1*t, can be used without any tendency to flaking of the layer or the like. As can be seen in Table 1, the different examples described here, do not show any tendency at all to any flaking, or the like.

All bend-tested samples passed the test with the ranking "accepted".

The six examples will now be described more in detail regarding their respective specifics:

5

Example 1

10 A 0,280 mm thick AISI 304 substrate material in a nominal composition of max 0,08% C, max 1% Si, max 2% Mn, 18.0-20.0% Cr, 8.0-10.0 % Ni and rest essentially Fe, and in annealed condition coated by using a strip speed of about 10 m/min with electron beam evaporation, with about 1
15 µm TiO directly on the substrate to get a bronze colour. The coated specimen with the TiO layer passed the bending test and also the scotch tape test. See sample 1 in Table 1 for the bending test result, scotch tape test and the measured *L**, *a** and *b** colour values.

20

Example 2

A 0,317 mm thick AISI 304 substrate material in a nominal composition of max 0,08% C, max 1% Si, max 2% Mn, 18.0-20.0% Cr, 8.0-10.0 % Ni and rest essentially Fe, and
25 in cold rolled condition coated using a strip speed of minimum 3 m/min by sputtering with TiN, TiAlN in several step to get multilayered structures. The mulitlayered coated material gets colours of blue, green and violet. The total layer thickness is 45 nm for the blue strip, 50 nm
30 for the purple strip and 100 nm for the green strip. The specimens with the coloured layers all passed the bending test and also the scotch tape tests. See samples 2-4 table 1 for the bending test result, scotch tape test and the measured *L**, *a** and *b** colour values.

Example 3

A 0,317 mm thick AISI 304 substrate material in a nominal composition of max 0,08% C, max 1% Si, max 2% Mn, 5 18.0-20.0% Cr, 8.0-10.0 % Ni and rest is essentially Fe, in cold rolled condition coated by using a strip speed of minimum 3 m/min by sputtering with TiAlN to get a black colour. The coated specimen with the black layer passed the bending test and also the scotch tape test. See sample 5 in 10 Table 1 for the bending test result, scotch tape test and for the measured L^* , a^* and b^* colour values.

Example 4

A 0,137 mm thick substrate material with the 15 designation Sandvik 13C26 with a nominal composition of 0,7 % C, 13 % Cr, 0,4 % Si and 0,7 % Mn (by weight) and rest essentially Fe, coated with TiN directly on to the steel by a sputtering process using a strip speed of minimum 3 m/min. The layer was coloured in gold. The sample was 20 afterwards heat-treated in 1100°C for 10s under N₂/H₂ atmosphere, which is a normal hardening procedure for this type of steel. See samples 6-7 in Table 1 for the bending test result, scotch tape test and for the measured L^* , a^* and b^* colour values. The difference in colour between the 25 gold coloured specimen before the heat-treatment and after the heat-treatment is measured to be $\Delta E = 6,5$.

Example 5

A 0,1 mm thick substrate material with the 30 designation Sandvik 13C26 with a nominal composition of 0,7 % C, 13 % Cr, 0,4 % Si and 0,7 % Mn (by weight) and rest essentially Fe coated using a strip speed of about 10 m/min by electron beam evaporation with about 0,1 µm titanium dioxide directly on the substrate to get a light turquoise

colour. The specimen with the TiO₂ layer passed the scotch tape test. See sample 8 in Table 1 for the bending test result, scotch tape test and for the measured L*, a* and b* colour values.

5

Example 6

A 0,1 mm thick substrate material with the designation Sandvik 13C26 with a nominal composition of 0,7 % C, 13 % Cr, 0,4 % Si and 0,7 % Mn (by weight) and rest 10 essentially Fe coated using a strip speed of minimum 3 m/min by electron beam evaporation with Cr₂O₃ directly on a previously TiN-coated substrate to get a black colour. The specimen with the Cr₂O₃ layer passed the bending test and also the scotch tape test. See sample 9 in Table 1 for the 15 bending test results, scotch tape test results and the measured L*, a* and b* colour values.

CLAIMS

1. A coated stainless steel strip product with an evenly distributed layer on one side or both sides of said strip characterized in that said layer has a decorative appearance, the thickness of said layer is maximally 10 μm ,
5 the tolerance of said layer is maximally +/- 30% of the layer thickness, the parameter value of L*, a*, b* is respectively $0 < L^* < 95$, $-66 < a^* < 64$, $-90 < b^* < 70$, the tolerance of said decorative appearance as expressed in
10 terms of ΔE is maximally 15 and that the layer has such a good adhesion so that the coated steel strip when tested in soft-annealed condition can be bent more than 90° over a radius maximally equal to $5*t$, where t is the thickness of
15 the said strip, without showing any tendency to flaking or the like.

2. Product according to claim 1 characterized in that the thickness of the strip substrate is between 0,015 mm and 3,0 mm

3. Product according to claim 1 or 2 characterized in
20 that the ratio between the thickness of the coating and the thickness of the strip is max 7%.

4. Product according to any of claims 1-3, characterized in that it is made of a substrate of ferritic stainless steel, austenitic stainless steel, stainless spring steel,
25 duplex stainless steel, hardenable chromium steel or precipitation hardenable stainless steel.

5. Product according to any of the preceding claims 1-4 characterized in that the substrate material in soft-annealed condition has a tensile strength of maximum 1400
30 MPa.

6. Product according to any of the claims 1-4 characterized in that the substrate material in cold-rolled condition has a tensile strength of minimum 500 MPa.

7. Product according to any of the claims 1-4 characterized in that the substrate material in hardened and/or tempered condition has a tensile strength of minimum 1000 MPa.

5 8. Product according to any of claims 1-7, characterized in that the coating is a binary metal oxide or a ternary metal oxide or mixtures or solid solutions of said binary metal oxides, the main ingredient in such a mixture or solid solution being Al_2O_3 , TiO_2 or Cr_2O_3 .

10 9. Product according to any of claims 1-7, characterized in that the coating is a coating of metal carbides or metal nitrides, preferably TiN , TiAlN , ZrN , TiC , or CrN , or mixtures thereof.

10. Product according to claim 9 subsequently heat-treated in a tempering or a hardening process using a suitable gas atmosphere, characterized in that the difference of the decorative appearance of the coating after the heat-treatment compared to before the heat-treatment as expressed in terms of ΔE is maximally 15.

20 11. Product according to claim 10 characterized in that the material after the subsequent heat-treatment has a tensile strength of more than 1000 MPa.

12. Product according to any of preceding claims, characterized in that the layer has a multi-layer constitution of up to 10 layers.

25 13. Product according to claim 12 characterized in that each individual layer has a thickness of between 0,01 to 10 μm .

14. Product according to claim 13, characterized in that 30 the layer has a multi-layer constitution of individual layers of different coatings of nitrides or carbides such as TiN and TiC , and if desired also in combination with layers of oxides in the form of Cr_2O_3 or Al_2O_3 or TiO_2 , or mixtures thereof.

15. Product according to claim 14, characterized in that there is also at least one covering layer of nickel or chromium or aluminum or titanium in thickness up to 2 μm .

16. Product according to any of claims 1-7 with a
5 desired decorative appearance achieved by using a two-step method with a coating applied in the first step and a subsequent processing done in the second step to achieve a desired colour, characterized in that the coating in the first step is a suitable covering layer of a metal such as
10 aluminum, chromium, titanium, zirconium or a binary oxide of said metal such as Al_2O_3 , TiO_2 , Cr_2O_3 , or mixtures of said metal and said binary oxide.

17. Product according to claim 16 characterized in that the material after the subsequent processing in the second-
15 step has a tensile strength of more than 1000 MPa.

18. Product according to any of claim 16-17 characterized in that the desired colour is achieved by incorporating a suitable element such as oxygen, carbon, nitrogen into the coating applied in the first step, by
20 using a reactive gas during a suitable heat-treatment.

19. Product according to any of claims 16-18 characterized in that the final product after the second step has a coating with a desired colour consisting of metal oxide, metal nitride, metal carbide, or a mixture such as
25 metal oxy-nitride, metal oxy-carbide or metal carbo-nitride.

20. Product according to any of claims 16-19 characterized in that the layer has a multi-layer constitution of up to 10 layers.

30 21. Product according to any of claims 16-20 characterized in that each individual layer has a thickness of between 0,01 to 10 μm .

22. Product according to any of the preceding claims 1-
21 characterized in a decorative appearance of a typical a

blue colour of normally $20 < L^* < 95$, $-66 < a^* < 64$ and $-83 < b^* < 0$.

23. Product according to any of the preceding claims
1-21 characterized in a decorative appearance of a typical
5 a green colour of normally $20 < L^* < 95$, $-66 < a^* < 0$ and $-83 < b^* < 70$.

24. Product according to any of the preceding claims
1-21 characterized in a decorative appearance of a
typical a red colour of normally $20 < L^* < 95$, $0 < a^* < 64$
10 and $-40 < b^* < 35$.

25. Product according to any of the preceding claims
1-21 characterized in a decorative appearance of a
typical golden colour of normally $20 < L^* < 95$, $-66 < a^* < 64$
and $0 < b^* < 70$.

15 26. Product according to any of the preceding
claims 1-21 characterized in a decorative appearance of a
typical black colour of $0 < L^* < 50$, $-20 < a^* < 20$ and $-20 < b^* < 20$.

20 27. Product according to any of the preceding
claims 1-21 characterized in a decorative appearance of a
typical violet colour of normally $20 < L^* < 95$, $20 < a^* < 60$
and $-25 < b^* < -60$.

25 28. A product according to any of the preceding claims
1-27, characterized in that it is suitable for cost
efficient and productive manufacturing of consumer
related applications, such as outdoor life applications,
sports and sea-life applications, household applications,
camera applications, mobile phones and other telecom
applications, edge applications such as knife, saw and
30 shaving applications or the like, and applications for
personal belongings and care such as watches, glasses,
cosmetic applications, buttons and zippers in clothing,
perfume bottles or the like.

29. Method of manufacturing a coated stainless steel strip product according to any of the preceding claims 1-
28, characterized in that said product is produced in a continuous roll-to-roll process with a minimum strip speed of 3 m/min, included in a strip production line using sputtering and/or electron beam evaporation comprising an etch chamber in-line.

1 / 5

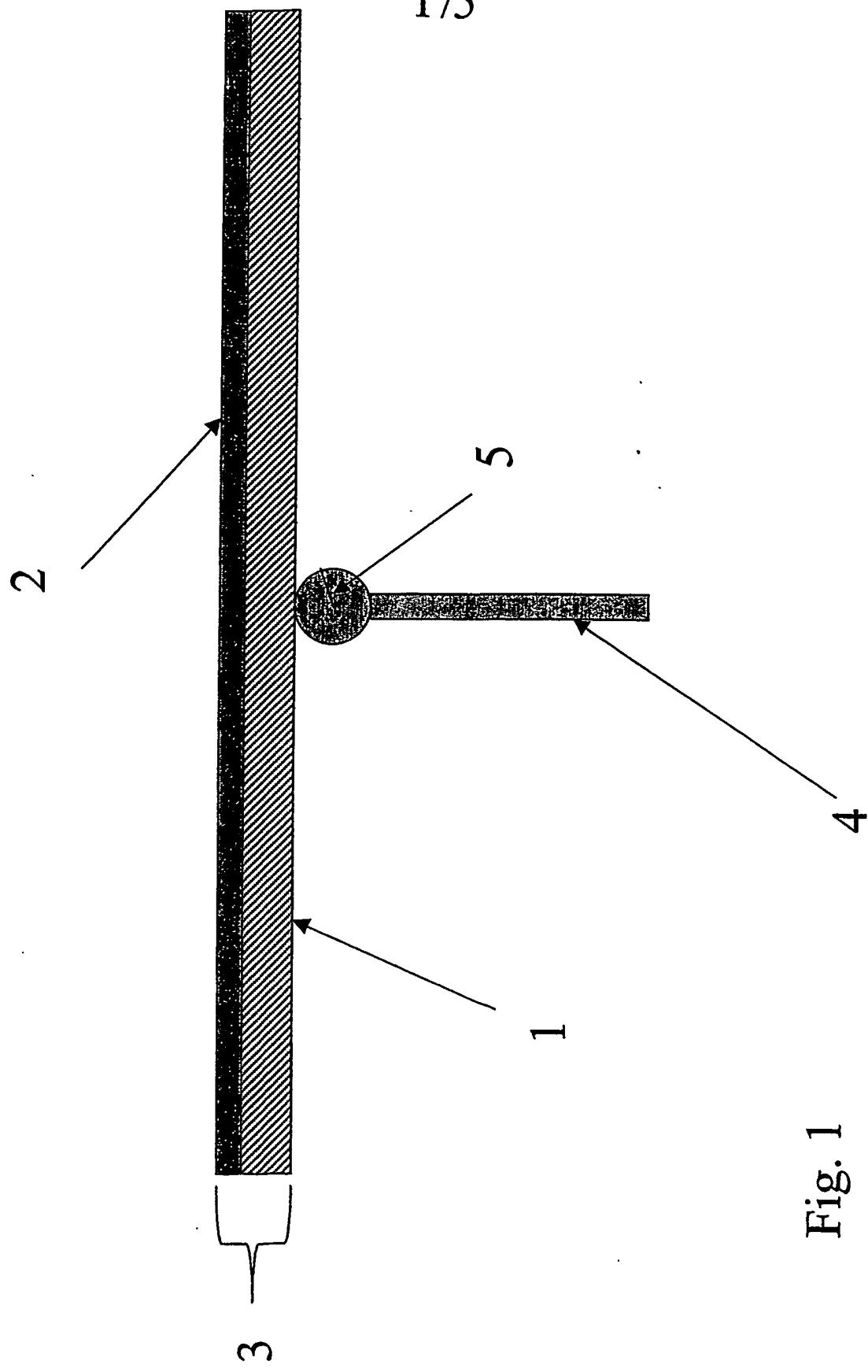


Fig. 1

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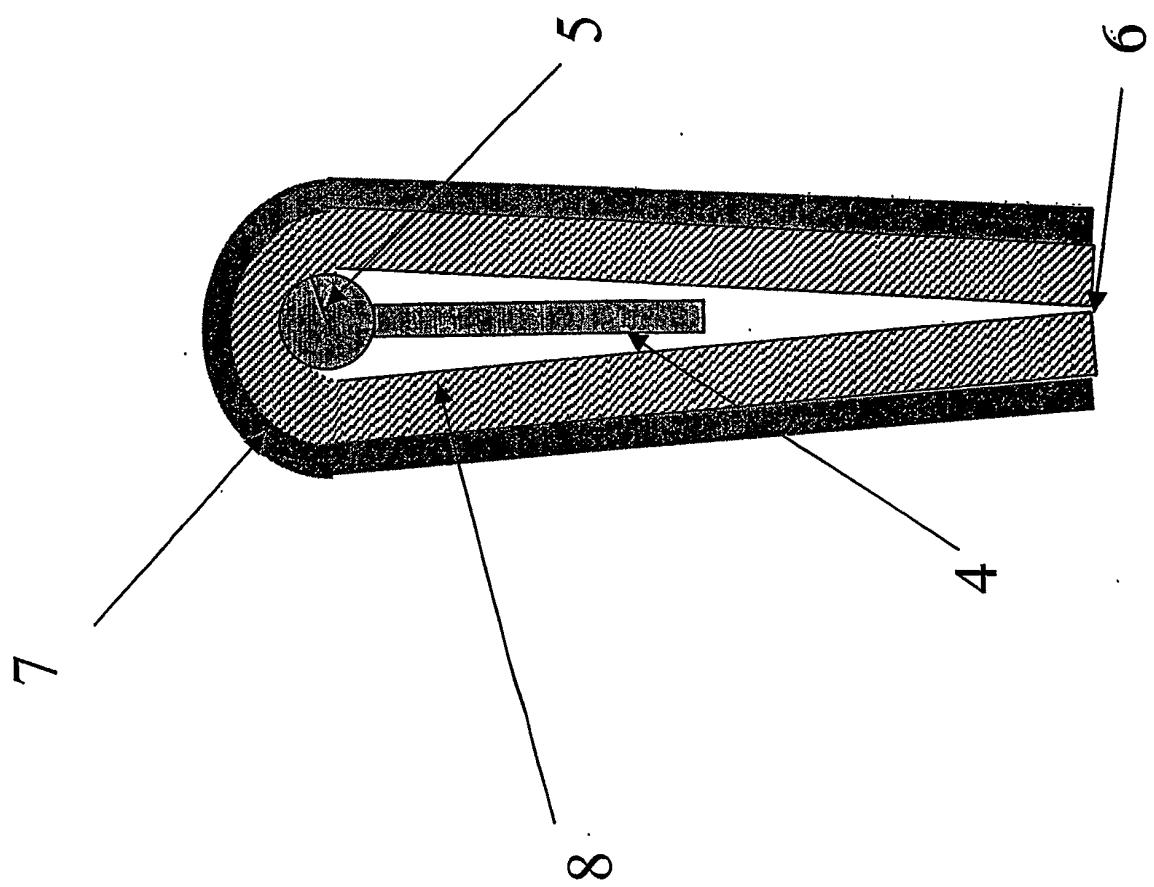


Fig. 2

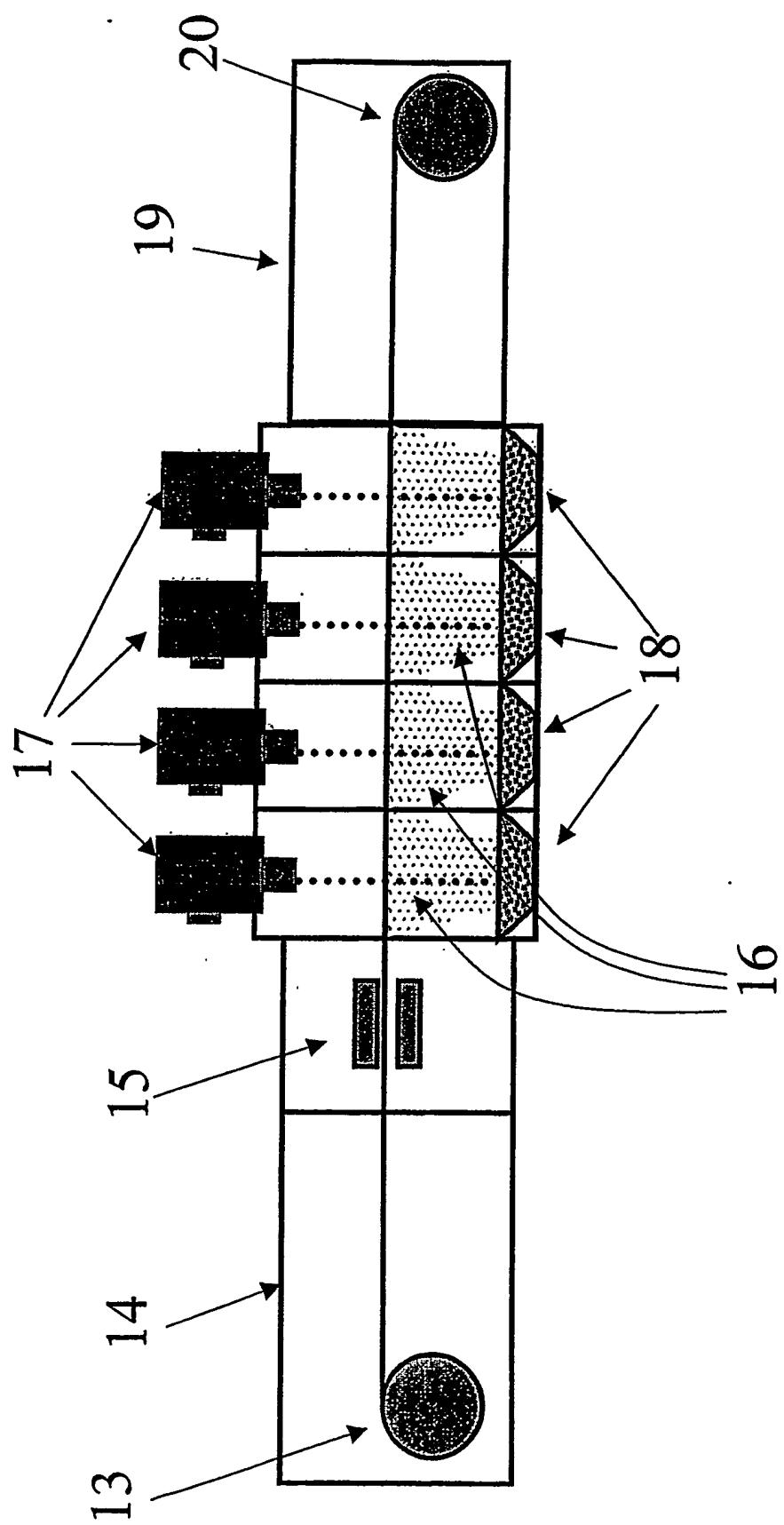


Fig. 3

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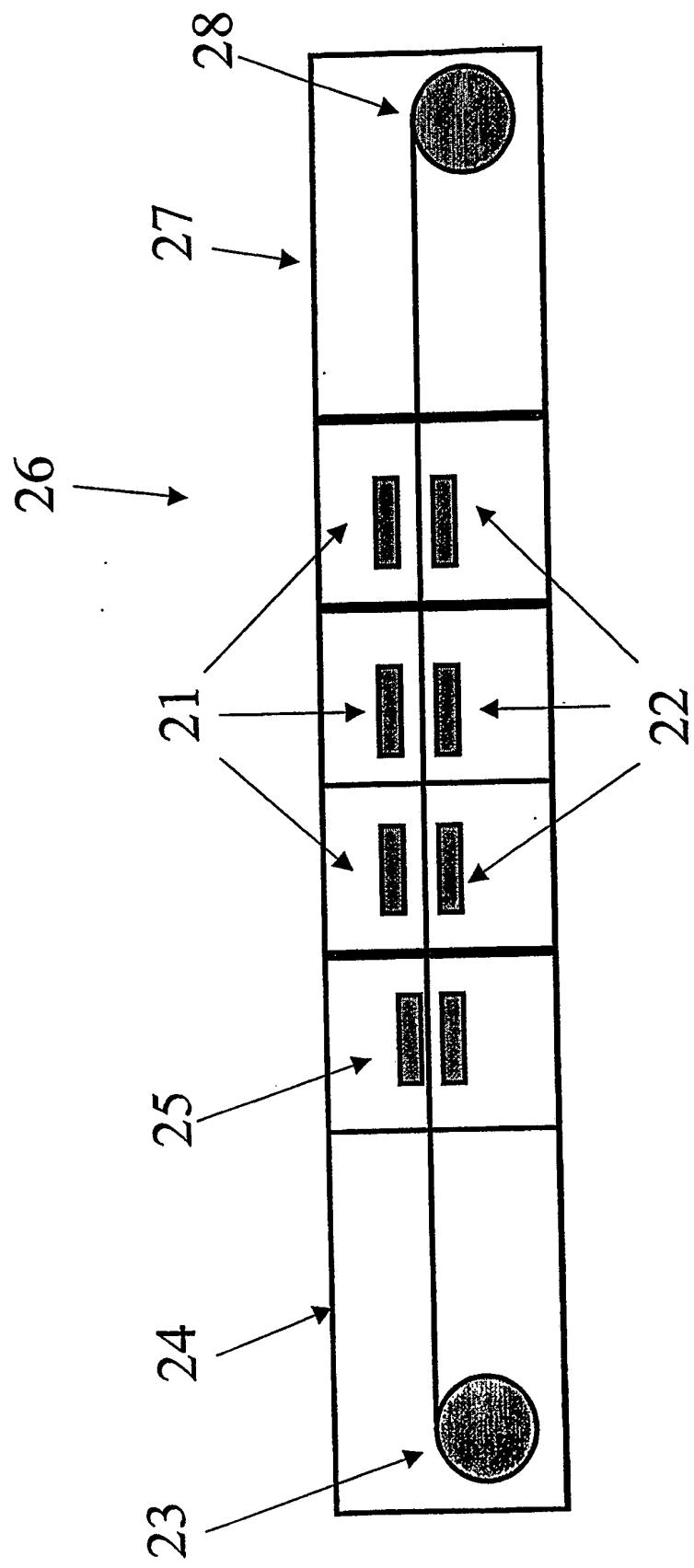


Fig. 4

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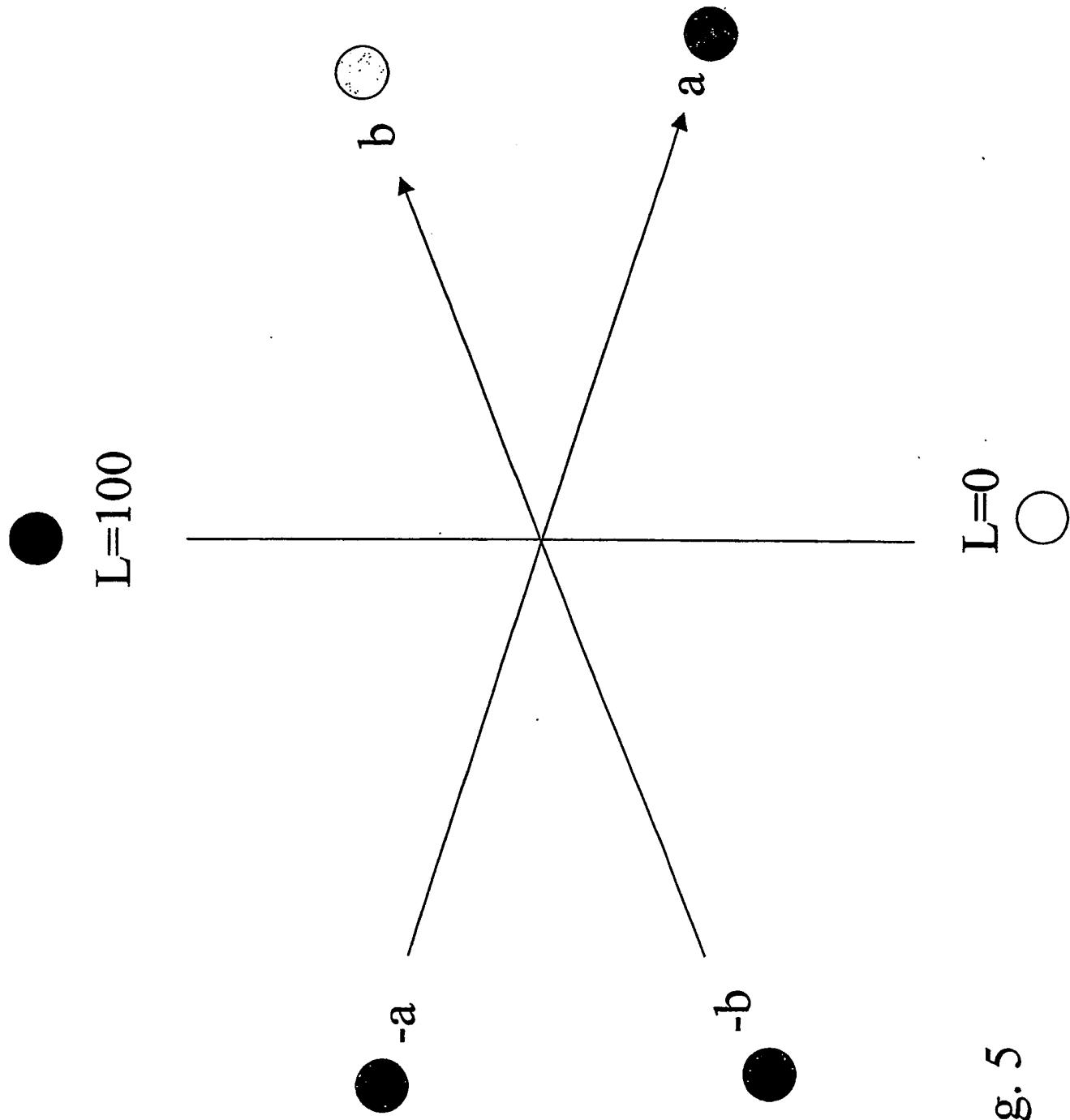


Fig. 5

INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 2004/001870

A. CLASSIFICATION OF SUBJECT MATTER

IPC7: C23C 14/56, C23C 14/30, C23C 14/34, C23C 14/06, C23C 14/08, C23C 14/16,
C23C 30/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC7: C23C

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE,DK,FI,NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

EOP-INTERNAL, WPI DATA

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 4763601 A (S. SAIDA ET AL), 16 August 1988 (16.08.1988), column 1, line 56 - line 64; column 3, line 18 - line 28; column 5, line 55 - line 57, column 7, line 46 - column 8, line 16, column 8, line 67 - column 10, line 11, figure 6 --	1-29
X	EP 0416887 A2 (NIPPON STEEL CORPORATION), 13 March 1991 (13.03.1991), page 2, line 1 - page 4, line 12; page 4, line 53 - page 6, line 10; page 6, line 36 - page 7, line 42 --	1-29
X	EP 0353060 A2 (NIPPON STEEL CORPORATION), 31 January 1990 (31.01.1990), page 3, line 7 - page 8, line 5 --	1-29

 Further documents are listed in the continuation of Box C. See patent family annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier application or patent but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

30 March 2005

Date of mailing of the international search report

06-04-2005

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INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 2004/001870

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	DATABASE WPI Week 199645 Derwent Publications Ltd., London, GB; Class M13, AN 1996-453225 & KR 9500308 B1 (KOREA RES ASSOC NEW IRON & STEEL MAKING), 13 January 1995 (1995-01-13) abstract --	1-29
X	US 4622919 A (Y. SUZUKI ET AL), 18 November 1986 (18.11.1986), column 2, line 54 - column 3, line 4; column 5, line 3 - column 7, line 37, figure 1 --	1-29
A	GB 1471977 A (JONES & LAUGHLIN STEEL CORPORATION), 27 April 1977 (27.04.1977), page 6, line 97 - line 101 --	29
X	DATABASE WPI Week 199249 Derwent Publications Ltd., London, GB; Class M13, AN 1992-403554 & JP 4301068 A (KOBE STEEL LTD), 23 October 1992 (1992-10-23) abstract --	1-29
X	PATENT ABSTRACTS OF JAPAN vol.017, no. 124 (C-1035), 16 March 1993 (1993-03-16) & JP 4301068 A (KOBE STEEL LTD), 23 October 1992 (1992-10-23) abstract --	1-29

INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 2004/001870

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	DATABASE WPI Week 199234 Derwent Publications Ltd., London, GB; Class M13, AN 1992-281782 & JP 4193968 A (KOBE STEEL LTD), 14 July 1992 (1992-07-14) abstract --	1-29
X	PATENT ABSTRACTS OF JAPAN vol. 016, no. 518 (C-0999), 26 October 1992 (1992-10-26) & JP 4193968 A (KOBE STEEL LTD), 14 July 1992 (1992-07-14) abstract --	1-29
A	US 5429843 A (K. MASAKI ET AL), 4 July 1995 (04.07.1995), column 3, line 29 ~ line 31 --	29
A	PATENT ABSTRACTS OF JAPAN vol. 014, no. 334 (C-0742), 18 July 1990 (1990-07-18) & JP 2122064 A (SUMITOMO METAL IND LTD), 09 May 1990 (1990-05-09) abstract --	12-15
E	WO 2005014876 A1 (SANDVIK AB), 17 February 2005 (17.02.2005), whole document --	1-29
E	WO 2005014877 A1 (SANDVIK AB), 17 February 2005 (17.02.2005), whole document -----	1-29

INTERNATIONAL SEARCH REPORTInternational application No.
PCT/SE2004/001870**Box No. II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)**

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

... / ...
2. Claims Nos.: 1 - 29 (partly)
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
... / ...
3. Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box No. III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.: .
4. No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.: .

Remark on Protest

- The additional search fees were accompanied by the applicant's protest.
 No protest accompanied the payment of additional search fees.

Box II

Present claims 1-28 relate to a product defined by reference to desirable characteristics or properties, namely:

- 1) parameters for the colour and tolerance of decorative appearance of a coating layer, i.e. L, a, b and ΔE,
- 2) thickness tolerance
- 3) adhesion of the coating to the steel strip
- 4) thickness of the layer

The use of all these desirable characteristics or properties in the present context is considered to lead to a lack of clarity within the meaning of Article 6 PCT. An attempt is made to define the product by reference to a result to be achieved. Furthermore, the claims relate to a large number of possible coated stainless steel strip products. Again, this lack of clarity render a meaningful complete search impossible.

Consequently, the search according to the invention in claims 1-29 has been restricted mainly to: coloured coated stainless strips coated with layers according to claims 8 and 9.

INTERNATIONAL SEARCH REPORT
Information on patent family members

01/03/2005

International application No.

PCT/SE 2004/001870

US	4763601	A	16/08/1988	DE	3786800 D,T 0305573 A,B	10/02/1994 08/03/1989
EP	0416887	A2	13/03/1991	DE	69018170 D,T JP 2619838 B JP 3097865 A US 5427843 A	27/07/1995 11/06/1997 23/04/1991 27/06/1995
EP	0353060	A2	31/01/1990	DE	68918196 D,T JP 2133565 A US 5079089 A US 5192410 A	16/02/1995 22/05/1990 07/01/1992 09/03/1993
US	4622919	A	18/11/1986	JP	60141869 A	26/07/1985
				US	4683149 A	28/07/1987
GB	1471977	A	27/04/1977	AU	481742 B AU 7329774 A CA 1026266 A CA 1050838 A DE 2446250 A FR 2245484 A,B IT 1022457 B JP 50061381 A	18/03/1976 18/03/1976 14/02/1978 20/03/1979 03/04/1975 25/04/1975 20/03/1978 26/05/1975
US	5429843	A	04/07/1995	DE	69220566 D,T EP 0572673 A,B JP 5320874 A JP 6052257 U WO 9310275 A KR 166098 B	19/02/1998 08/12/1993 07/12/1993 15/07/1994 27/05/1993 15/01/1999
WO	2005014876	A1	17/02/2005	SE	0302207 D	00/00/0000
				WO	2005014877 A	17/02/2005
WO	2005014877	A1	17/02/2005	SE	0302207 D	00/00/0000
				WO	2005014876 A	17/02/2005

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